(FILE 'HOME' ENTERED AT 12:54:37 ON 22 AUG 2003)

	FILE	'REGISTRY'	ENTERED	ΑT	12:54:5	O ON	22	AUG	2003				
L1		STRUG	CTURE UP	LOA	DED								
L2		0 S L1	SSS SAM										
L3		0 S L1	SSS FUL	L									
L4		STRUC	CTURE UP	LOA	DED								
L5		0 S L4	SSS SAM										
L6		0 S L4	SSS FUL	L									
L7		STRU	CTURE UP	LOA:	DED								
L8		0 S L7	SSS SAM										
L9		9 S L 7	SSS FUL	L									
	FILE	'CAPLUS, M	EDLINE,	USP.	ATFULL'	ENTE	RED	ΑT	13:01:27	ON	22	AUG	2003
1.10		4 S L9											

L10 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2003:3

2003:356095 CAPLUS

DOCUMENT NUMBER:

138:338411

TITLE:

Preparation of oligonucleotide labeling reactants based on acyclic nucleosides and conjugates derived

thereof

INVENTOR(S):

Hovinen, Jari Wallac Oy, Finland

PATENT ASSIGNEE(S): SOURCE:

Eur. Pat. Appl., 31 pp.

CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1308452	A2	20030507	EP 2002-396153	20021010

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,

IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK
US 2003118999 A1 20030626 US 2001-985454 20011102
PRIORITY APPLN. INFO.: US 2001-985454 A 20011102

IT 518027-22-0P 518027-23-1P

RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (prepn. of oligonucleotide labeling reactants based on acyclic nucleosides and conjugates derived thereof)

RN 518027-22-0 CAPLUS

CN Glycine, N,N'-[[4'-[4-[6-[3-[(2S)-3-[bis(4-methoxyphenyl)phenylmethoxy]-2-hydroxypropyl]-3,6-dihydro-2,6-dioxo-1(2H)-pyrimidinyl]-1-hexynyl]phenyl][2,2':6',2''-terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

RN 518027-23-1 CAPLUS
CN Glycine, N,N'-[[4'-[4-[6-[3-[(2S)-3-[bis(4-methoxyphenyl)phenylmethoxy]-2[[[bis(1-methylethyl)amino](2-cyanoethoxy)phosphino]oxy]propyl]-3,6dihydro-2,6-dioxo-1(2H)-pyrimidinyl]-1-hexynyl]phenyl][2,2':6',2''terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-,
dimethyl ester (9CI) (CA INDEX NAME)

L10 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2003 ACS on STN

2001:490069 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER:

135:242452

Versatile Strategy for Oligonucleotide Derivatization. TITLE:

Introduction of Lanthanide(III) Chelates to

Oligonucleotides

Hovinen, Jari; Hakala, Harri AUTHOR(S):

PerkinElmer Life Sciences Wallac Oy, Turku, FIN-20101, CORPORATE SOURCE:

Finland

Organic Letters (2001), 3(16), 2473-2476 SOURCE:

CODEN: ORLEF7; ISSN: 1523-7060

American Chemical Society PUBLISHER:

DOCUMENT TYPE: Journal English LANGUAGE:

CASREACT 135:242452 OTHER SOURCE(S):

358978-79-7P 358978-80-0P 358978-84-4P

358978-85-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(versatile strategy for oligonucleotide derivatization introduction of lanthanide chelates to oligonucleotides)

RN 358978-79-7 CAPLUS

Glycine, N,N'-[[4'-[4-(6-hydroxy-1-hexynyl)phenyl][2,2':6',2''-terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-, CN dimethyl ester (9CI) (CA INDEX NAME)

$$\begin{array}{c|c}
O & CH_2-C-OMe \\
MeO-C-CH_2-N-CH_2 & O & CH_2-C-OMe \\
N & O & CH_2-C-OMe \\
N & O & CH_2-C-OMe \\
CH_2-N-CH_2-C-OMe & O & CH_2-C-OMe \\
CH_2-N-CH_2-C-OMe & O & O & O \\
\end{array}$$

RN 358978-80-0 CAPLUS
CN Glycine, N,N'-[[4'-[4-[6-[3-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-2-deoxy-.beta.-D-erythro-pentofuranosyl]-3,6-dihydro-2,6-dioxo-1(2H)-pyrimidinyl]-1-hexynyl]phenyl][2,2':6',2''-terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

PAGE 1-A

Absolute stereochemistry.

MeO

HO

RN 358978-84-4 CAPLUS

Glycine, N,N'-[[4'-[4-[6-[3-[5-0-[bis(4-methoxyphenyl)phenylmethyl]-3-0-[bis(1-methylethyl)amino](2-cyanoethoxy)phosphino]-2-deoxy-.beta.-D-erythro-pentofuranosyl]-3,6-dihydro-2,6-dioxo-1(2H)-pyrimidinyl]-1-hexynyl]phenyl][2,2':6',2''-terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

RN 358978-85-5 CAPLUS
CN Glycine, N,N'-[[4'-[4-[6-[[[bis(1-methylethyl)amino](2-cyanoethoxy)phosphino]oxy]-1-hexynyl]phenyl][2,2':6',2''-terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

NC-CH₂-CH₂-O-P-O-(CH₂)₄-C=C

```
C-OMe
REFERENCE COUNT:
                         12
                               THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L10 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2003 ACS on STN
                         2000:553830 CAPLUS
ACCESSION NUMBER:
                         133:304860
DOCUMENT NUMBER:
                         Synthesis and spectral properties of a new luminescent
TITLE:
                         europium(III) terpyridyl chelate
                         Cooper, Michael E.; Sammes, Peter G.
AUTHOR(S):
                         Amersham Pharmacia Biotech, Whitchurch, Cardiff, CF14
CORPORATE SOURCE:
                         7YT, UK
                         Perkin 2 (2000), (8), 1695-1700
SOURCE:
                         CODEN: PRKTFO
                         Royal Society of Chemistry
PUBLISHER:
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     300775-34-2P 300775-36-4P 300775-38-6P
IΤ
    RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (intermediate product in prepn. of luminescent europium
        terpyridylbis (methylamine) tetraacetate complex)
RN
     300775-34-2 CAPLUS
     Glycine, N, N'-[[4'-(4-methylphenyl)[2,2':6',2''-terpyridine]-6,6''-
CN
```

divl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-, dimethyl ester (9CI)

--- OMe

(CA INDEX NAME)

O
$$CH_2-C-OMe$$

MeO-C-CH₂-N-CH₂

N

CH₂-C-OMe

CH₂-C-OMe

CH₂-C-OMe

RN 300775-36-4 CAPLUS
CN Glycine, N,N'-[[4'-(4-methyl-3-nitrophenyl)[2,2':6',2''-terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

RN 300775-38-6 CAPLUS
CN Glycine, N,N'-[[4'-(3-amino-4-methylphenyl)[2,2':6',2''-terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

O
$$CH_2-C-OMe$$

MeO-C- CH_2-N-CH_2

NH2

MeO-C- $CH_2-N-CH_2-C-OMe$

CH2-N- $CH_2-C-OMe$

40

L10 ANSWER 4 OF 4 USPATFULL on STN

ACCESSION NUMBER: 2003:173155 USPATFULL

TITLE: Oligonucleotide labeling reactants based on

acyclonucleosides and conjugates derived thereof

INVENTOR(S): Hovinen, Jari, Raisio, FINLAND

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: James C. Lydon, Attorney at Law, Suite 100, 100

Daingerfield Road, Alexandria, VA, 22314

NUMBER OF CLAIMS: 17 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 1 Drawing Page(s)

LINE COUNT: 871

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 518027-22-0P 518027-23-1P

(prepn. of oligonucleotide labeling reactants based on acyclic

nucleosides and conjugates derived thereof)

RN 518027-22-0 USPATFULL

CN Glycine, N, N' - [[4' - [4 - [6 - [3 - [(2S) - 3 - [bis(4 - methoxyphenyl)phenylmethoxy] - 2 - [bis(4 - methoxyphenyl)phenylmethoxy]]

hydroxypropyl]-3,6-dihydro-2,6-dioxo-1(2H)-pyrimidinyl]-1-

hexynyl]phenyl][2,2':6',2''-terpyridine]-6,6''-diyl]bis(methylene)]bis[N-

(2-methoxy-2-oxoethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

RN 518027-23-1 USPATFULL

Glycine, N,N'-[[4'-[4-[6-[3-[(2S)-3-[bis(4-methoxyphenyl)phenylmethoxy]-2[[[bis(1-methylethyl)amino](2-cyanoethoxy)phosphino]oxy]propyl]-3,6dihydro-2,6-dioxo-1(2H)-pyrimidinyl]-1-hexynyl]phenyl][2,2':6',2''terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-,
dimethyl ester (9CI) (CA INDEX NAME)

=>

=> d 11

L1 HAS NO ANSWERS

L1 STR

Structure attributes must be viewed using STN Express query preparation.

=> s 11 sss sam

SAMPLE SEARCH INITIATED 14:22:33 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED - 134 TO ITERATE

0 ANSWERS 134 ITERATIONS 100.0% PROCESSED

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 1986 TO 3374

0 TO PROJECTED ANSWERS:

L2 0 SEA SSS SAM L1

=> s 11 sss full

FULL SEARCH INITIATED 14:22:41 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 3041 TO ITERATE

3041 ITERATIONS 0 ANSWERS 100.0% PROCESSED

SEARCH TIME: 00.00.01

L3 O SEA SSS FUL L1

=> Uploading 09847384-7.str

STRUCTURE UPLOADED L4

Structure attributes must be viewed using STN Express query preparation.

=> s 14 sss sam

SAMPLE SEARCH INITIATED 14:24:14 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 2 TO ITERATE

100.0% PROCESSED 2 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**

PROJECTED ITERATIONS: 2 TO 124 PROJECTED ANSWERS: 0 TO 0

L5 0 SEA SSS SAM L4

=> s 14 sss full

FULL SEARCH INITIATED 14:24:21 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 67 TO ITERATE

100.0% PROCESSED 67 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

L6 0 SEA SSS FUL L4

=> Uploading 09847384-8.str

L7 STRUCTURE UPLOADED

=> d 17

L7 HAS NO ANSWERS

Structure attributes must be viewed using STN Express query preparation.

=> s 17 sss sam SAMPLE SEARCH INITIATED 14:27:21 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED - 2 TO ITERATE

100.0% PROCESSED 2 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
PROJECTED ITERATIONS: 2 TO 124
PROJECTED ANSWERS: 0 TO 0

PROJECTED ANSWERS: U TO

L8 0 SEA SSS SAM L7

=> s 17 sss full FULL SEARCH INITIATED 14:27:27 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 82 TO ITERATE

100.0% PROCESSED 82 ITERATIONS 20 ANSWERS

SEARCH TIME: 00.00.01

L9 20 SEA SSS FUL L7

=> file caplus medline uspatfull

COST IN U.S. DOLLARS

SINCE FILE
ENTRY
SESSION
FULL ESTIMATED COST

447.25

447.46

FILE 'CAPLUS' ENTERED AT 14:28:00 ON 22 AUG 2003 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'MEDLINE' ENTERED AT 14:28:00 ON 22 AUG 2003

FILE 'USPATFULL' ENTERED AT 14:28:00 ON 22 AUG 2003 CA INDEXING COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS)

=> s 19

L10 16 L9

=> s 110 and label

0 L10 AND LABEL L11

 \Rightarrow s 110 and DMTr

L120 L10 AND DMTR

=> s 110 and dimethoxytrityl

3 L10 AND DIMETHOXYTRITYL L13

=> d 113 1-3 ibib abs hitstr

L13 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

2003:227376 CAPLUS

DOCUMENT NUMBER:

138:255457

TITLE:

Thymidine derivatives having an intercalator at N3

position of thymine base and use in antisense

oligonucleotides

INVENTOR(S):

Ono, Akira

PATENT ASSIGNEE(S):

Yamasa Shoyu Co., Ltd., Japan Jpn. Kokai Tokkyo Koho, 8 pp.

SOURCE:

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

1

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2003088374	A2	20030325	JP 2001-279240	20010914
PRIORITY APPLN. INFO.	:		JP 2001-279240	20010914
GI				

Thymidine derivs. having an intercalator introduced at N3 position of AB thymine base represented by (I) (A = intercalator, B = linker, X = H or -OY where Y = lower alkyl group which may be substituted, R1 and R2 = H, protective group or substituted phosphino), and oligonucleotides contg. them, are disclosed. The presence of 3'-0-[(2cyanoethoxy) (diisopropylamino) phosphino] -5'-0-dimethoxytrityl -N3-[2- (anthraquinone-2-carboxyamino)ethyl] thymidine (chem. compd. 1:ant) ODNs was found to effectively stabilize duplex and triplex formation.

501912-34-1P ΙT

RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(thymidine derivs. having an intercalator at N3 position of thymine base and use in antisense oligonucleotides)

RN 501912-34-1 CAPLUS

CN

Thymidine, 5'-O-[bis(4-methoxyphenyl)phenylmethyl]-3-[2-[[(9,10-dihydro-9,10-dioxo-2-anthracenyl)carbonyl]amino]ethyl]-, 3'-[2-cyanoethylbis(1-methylethyl)phosphoramidite] (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-B

^{_}OMe

L13 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2001:490069 CAPLUS

DOCUMENT NUMBER: 135:242452

TITLE: Versatile Strategy for Oligonucleotide Derivatization.

Introduction of Lanthanide(III) Chelates to

Oligonucleotides

AUTHOR(S): Hovinen, Jari; Hakala, Harri

CORPORATE SOURCE: PerkinElmer Life Sciences Wallac Oy, Turku, FIN-20101,

Finland

SOURCE: Organic Letters (2001), 3(16), 2473-2476

CODEN: ORLEF7; ISSN: 1523-7060

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 135:242452

AB Novel nucleosidic phosphoramidite blocks were synthesized by a Mitsunobu reaction between 2'-deoxy-5'-O-(4,4'-dimethoxytrityl) uridine and a primary alc. contg. a conjugate group in its structure (a protected functional group, an org. dye, or a precursor of a lanthanide(III) chelate) followed by phosphitylation. They were used in machine-assisted DNA synthesis in the std. manner. A slightly modified deprotection procedure was used for the prepn. of oligonucleotide conjugates tethered to lanthanide(III) chelates. For the latter application one non-nucleosidic block was also synthesized.

IT 358978-81-1P 358978-82-2P 358978-83-3P 358978-84-4P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(versatile strategy for oligonucleotide derivatization introduction of lanthanide chelates to oligonucleotides)

RN 358978-81-1 CAPLUS

CN Uridine, 5'-0-[bis(4-methoxyphenyl)phenylmethyl]-2'-deoxy-3-[6-[(trifluoroacetyl)amino]hexyl]-, 3'-[2-cyanoethyl bis(1-methylethyl)phosphoramidite] (9CI) (CA INDEX NAME)

Absolute stereochemistry.

$$F_{3}C$$

N

(i-Pr) 2N

P

CN

OMe

OMe

RN 358978-82-2 CAPLUS

CN Glycine, N, N'-[[4-[6-[3-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-3-O-[bis(1-methylethyl)amino](2-cyanoethoxy)phosphino]-2-deoxy-.beta.-D-erythro-pentofuranosyl]-3,6-dihydro-2,6-dioxo-1(2H)-pyrimidinyl]-1-hexynyl]-2,6-pyridinediyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-,dimethyl ester (9CI) (CA INDEX NAME)

MeO
$$C = C - (CH_2) \frac{1}{4}$$
 $C = C - (CH_2) \frac{1}{4}$ $C = C - (CH_2)$

PAGE 1-B

RN 358978-83-3 CAPLUS

CN Uridine, 5'-O-[bis(4-methoxyphenyl)phenylmethyl]-2'-deoxy-3-[6-[[[4-[(1E)-[4-(dimethylamino)phenyl]azo]phenyl]sulfonyl]amino]hexyl]-,
3'-[2-cyanoethyl bis(1-methylethyl)phosphoramidite] (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

PAGE 1-B

RN 358978-84-4 CAPLUS

Glycine, N,N'-[[4'-[4-[6-[3-[5-0-[bis(4-methoxyphenyl)phenylmethyl]-3-0-[bis(1-methylethyl)amino](2-cyanoethoxy)phosphino]-2-deoxy-.beta.-D-erythro-pentofuranosyl]-3,6-dihydro-2,6-dioxo-1(2H)-pyrimidinyl]-1-hexynyl]phenyl][2,2':6',2''-terpyridine]-6,6''-diyl]bis(methylene)]bis[N-(2-methoxy-2-oxoethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 1999:559569 CAPLUS

DOCUMENT NUMBER: 131:257798

TITLE: Synthesis of oligodeoxynucleotides containing

5-(.beta.-D-glucopyranosyloxymethyl)-2'-deoxyuridine, a modified nucleoside in the DNA of Trypanosoma brucei

AUTHOR(S): De Kort, Martin; Ebrahimi, Edwin; Wijsman, Eric R.;

Van der Marel, Gijs A.; Van Boom, Jacques H.

CORPORATE SOURCE: Leiden Institute Chemistry, Gorlaeus Laboratories,

Univ. Leiden, Leiden, 2300 RA, Neth.

SOURCE: European Journal of Organic Chemistry (1999), (9),

2337-2344

CODEN: EJOCFK; ISSN: 1434-193X

PUBLISHER: Wiley-VCH Verlag GmbH

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 131:257798

GI

The synthesis of the recently discovered modified DNA base 5-(.beta.-D-glucopyranosyloxymethyl)-2'-deoxyuridine (.beta.-dJ) is described. Me3SiO3SCF3-mediated .beta.-glucosylation of a 5-(hydroxymethyl)-2'-deoxyuridine (5-HMdU) deriv., obtained in 20% yield from 2'-deoxyuridine, with 2,3,4,6-tetra-O-benzoyl-.alpha.-D-glucopyranosyl trichloroacetimidate (I) gave dimer II [R = H; R1R2 = Si(CHMe2)2OSi(CHMe2)2] in 47% yield. On the other hand, condensation of I with a N3-(pivaloyloxymethyl)-protected deriv., readily available from thymidine in 48% yield, afforded the fully protected nucleoside II (R = CH2O2CMe3; R1, R2 = SiMe2CMe3) in 96% yield. The latter compd. was converted into phosphoramidite II {R = CH2O2CMe3, R1 = 4,4'-dimethoxytrityl, R2 = P[O(CH2)2CN]N(CHMe2)2}, which was applied in the automated solid-phase synthesis of several biol. interesting .beta.-dJ-contg. DNA fragments.

IT 244631-58-1P

RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of (glucopyranosyloxymethyl)deoxyuridine-contg. oligodeoxynucleotides)

RN 244631-58-1 CAPLUS

CN Thymidine, 5'-O-[bis(4-methoxyphenyl)phenylmethyl]-3-[(2,2-dimethyl-1-oxopropoxy)methyl]-.alpha.-[(2,3,4,6-tetra-O-benzoyl-.beta.-D-glucopyranosyl)oxy]-, 3'-[2-cyanoethyl bis(1-methylethyl)phosphoramidite]
(9CI) (CA INDEX NAME)

Absolute stereochemistry.

36

REFERENCE COUNT:

THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS on STN
L1
     14470-28-1 REGISTRY
RN
     Benzene, 1-(chlorodiphenylmethyl)-4-methoxy- (9CI)
                                                         (CA INDEX NAME)
CN
OTHER CA INDEX NAMES:
    Anisole, p-(chlorodiphenylmethyl)- (6CI, 7CI, 8CI)
CN
OTHER NAMES:
     (p-Anisyl)diphenylmethyl chloride
CN
     1-(Chlorodiphenylmethyl)-4-methoxybenzene
CN
     4-Anisyl (chloro) diphenylmethane
CN
     4-Methoxytriphenylmethyl chloride
CN
CN
     4-Methoxytrityl chloride
CN
     4-Monomethoxytrityl chloride
     Mono-p-methoxytrityl chloride
CN
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CN
CN
     p-(Chlorodiphenylmethyl)anisole
CN
     p-Anisylchlorodiphenylmethane
CN
     p-Methoxytrityl chloride
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MF
     STN Files:
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LC
       CHEMINFORMRX, CHEMLIST, CSCHEM, IFICDB, IFIPAT, IFIUDB, SYNTHLINE,
       TOXCENTER, USPAT2, USPATFULL
         (*File contains numerically searchable property data)
     Other Sources:
                      EINECS**
         (**Enter CHEMLIST File for up-to-date regulatory information)
```

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

273 REFERENCES IN FILE CA (1937 TO DATE)

8 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

273 REFERENCES IN FILE CAPLUS (1937 TO DATE)

2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)